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- (54) SILICONE MODIFIED PHOSPHOLIPID COMPOSITIONS

 SILIKON-MODIFIZIERTE PHOSPHOLIPID-ZUSAMMENSETZUNGEN

 COMPOSITIONS DE PHOSPHOLIPIDES MODIFIES AU SILICONE
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Description

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[0001] The present invention relates to novel organosilicone compounds according to the preamble of claims 1 and 5 and, more particularly, to silicone containing derivatives having at least one esterified phosphate group in the molecule and it relates also to a method of preparing novel phospholipid compositions according to the preamble of claim 10. [0002] Phosphate esters, quaternary amine compounds, betaines and certain substituted betaines are known in the art and have been commercially used over the years for a variety of applications, including those requiring surfactant properties. More recently, various betaine derivatives having, in general, specific quaternary compounds linked to phosphate esters referred to as phosphobetaines, and more particularly "synthetic phospholipids," have been disclosed, for example, in U.S.-A-4,215,064,4,233,192,4,380,637 and 4,302,036 to Lindemann et al.; U.S. -A-4,209,449,4,336,385 and 4,503,002 to Mayhew et al.; U.S. -A-4,243,602,4,283,542 and 4,336,386 to O'Lenick et al; and U.S. -A-4,617,404 to Lukenbach et al. These synthetic phospholipids are disclosed as exhibiting outstanding foaming, viscosity building, wetting, cleansing, detergency, anti-static, conditioning and emulsifying properties, making them useful in industrial applications calling for high performance surface active agents. The synthetic phospholipids are also described as being highly stable compositions which are well tolerated by human tissue (i.e. they exhibit exceptionally low oral toxicity and ocular irritation) and, hence, are well suited for use in a variety of personal care applications including cosmetic formulations as well as in industrial processes.

[0003] A variety of organosiloxane compositions including compositions which exhibit excellent properties as surface active agents, lubricants and the like are well known and have been used commercially over the years, including for personal care and home care applications. In general, organosiloxane compositions are water-insoluble and the costs thereof are greater than many other commercial materials which has limited their use for many applications. Recently, particular types of betaine and phosphobetaine modified organosiloxanes have been disclosed, for example, in U.S. -A- 4,609,750 and 4,654,161 to Kollmeier et al. and U.S. -A- 5,091,493 to O'Lenick et al which have been suggested as exhibiting high foaming characteristics in water, substantivity to a variety of surfaces, reduced irritation to the eyes and skin and improved, although limited, water-solubility properties. While, as indicated, certain organosilicone compositions containing phosphobetaines and methods for preparing the same have been suggested, there has been no prior disclosure or suggestion of the novel silicone modified phospholipid compositions and preparation methods or of the novel silicone modified phospholipid compositions and methods for preparing the same herein described which compositions exhibit a greater range of properties for different applications as well as providing means for closer control of the costs of the organosilicone products. The phospholipid compositions are described in the characterizing portion of claims 1 and 5 and the method of preparing novel phospholipid compositions is disclosed in the characterizing portion of claims 10.

[0004] It is accordingly an object of the present invention to provide novel silicone-modified phospholipid compositions suitable for use in solvent and/or preferably aqueous based systems which exhibit excellent surface-active properties including high foaming, are well tolerated by human tissue, are substantive to the surface of natural and synthetic fiber, and the like.

[0005] It is another object of the present invention to provide novel water-soluble silicone-modified phospholipid compositions containing terminal, lateral (pendant) or combinations of terminal and lateral (pendant) silicone moieties and wherein the compositions can be prepared with a variety of concentrations of silicone as desired or required.

[0006] In accordance with the present invention, there has now been discovered novel phospholipid compositions that may be represented by the following general formula:

$$\begin{array}{c}
O \\
R - Y - O - P \\
A_1
\end{array}$$

wherein:

A is selected from H, M, and R-Y-;
A₁ is selected from H, OH, OM, and R-Y-O-;
x is o or an interger from 1 to 5;
M is a cation, preferably an alkali metal;
Y is alkylene or substitued alkylene; and
R is selected from;

a) a quaternized organosilicone amidoamine moiety of the formula:

$$R_{1} = \frac{\begin{pmatrix} CII_{2} \end{pmatrix}_{n1} B_{n2}}{\begin{pmatrix} CII_{2} \end{pmatrix}_{n3}} \begin{pmatrix} CII_{2} \end{pmatrix}_{n3} \begin{pmatrix} CII_{2} \end{pmatrix}_{n} \begin{pmatrix} CII_$$

wherein:

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R₁ is a silicone backbone chain as hereinafter described to which amidoamine and/or amine functional

group(s) as herein described can be attached;

R₂ is hydrogen or alkyl, hydroxyalkyl or alkenyl of up to 6 carbon atoms each, or cycloalkyl of up to 6 carbon atoms, or polyoxyalkylene of up to 10 carbon atoms preferably from 2 to 5 carbon atoms

within the oxyalkylene unit;

R₃ and R₄, which may be the same or different, are selected from alkyl, hydroxyalkyl, carboxyalkyl of up to 6 carbon atoms in each alkyl, and polyoxyalkylene of up to 10 carbon atoms; in addition R, and R₄

taken together with the nitrogen to which they are attached represent a N-heterocycle;

X is an anion, preferably a halogen;

n is an integer from 2 to 12;

n1 is zero or an integer from 1 to 12;

n² is 0 or 1;

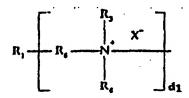
n³ is an integer from 1 to 5;

B is sulfur (S) or oxygen (O); with the proviso that when n' is 0, n¹ or n³ is at least 1 and when

n' is 1, n' and n' each is at least 1; and

d is one or greater, preferably 2-10;

b) a quaternized organosilicone tertiary amine moiety of the formula:



wherein:

 R_6

R₁ is a silicone backbone chain as hereinafter described to which amidoamine and/or amine functional groups can be attached;

is alkylene, hydroxyalkylene, arylene, alkarylene, aralkylene, heteroalkylene wherein the heteroatom

can be N, S or O and there can be more than one of such hetero atoms in the chain;

X- is an anion, preferably a halogen;

d¹ is an integer of one or greater, preferably from 2 to 10; and

R₃ and R₄ are as hereinabove defined;

or a mixture of the quaternized organosilicone amidoamine moiety "a" and/or quaternized tertiary amine moiety "b" above and the moieties:

c) a quaternized organic amidoamine moiety of the formula:

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$$\begin{bmatrix} O & \begin{bmatrix} R_2 & O \\ \\ R_3 - C & \end{bmatrix} & \begin{bmatrix} R_2 & R_3 \\ \\ N & C \end{bmatrix} & \begin{bmatrix} R_2 & R_3 \\ \\ R_4 & \end{bmatrix} & \begin{bmatrix} R_3 & C \\ \\ R_4 & \end{bmatrix}$$

wherein:

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R₅ is alkyl, alkenyl, alkoxyalkyl or hydroxyalkyl of from 5 to 21 carbon atoms each, alkaryl or aryl of up

to 20 carbon atoms;

R₂ is hydrogen or alkyl, hydroxyalkyl or alkenyl of up to 6 carbon atoms each, cycloalkyl of up to 6 carbon

atoms, or polyoxyalkylene of up to 10 carbon atoms, preferably of from 2 to 5 carbon atoms, within

the oxyalkylene unit;

R₃ and R₄ which may be the same or different, are selected from alkyl, hydroxyalkyl, carboxyalkyl of up to 6

carbon atoms in each alkyl moiety, and polyoxyalkylene of up to 10 carbon atoms; in addition R₃ and R₄ taken together with the nitrogen to which they are attached may represent an N-heterocycle;

X is an anion, preferably a halogen;

g is 0 or 1;

n is integer from 2 to 12; and

n⁴ is 1 or greater; and/or

d) an organic quaternized tertiary amine moiety of the formula:

$$\begin{bmatrix} R_{13} & X^{-} \\ \vdots & X^{-} \\ \vdots & \vdots \\ R_{15} \end{bmatrix}$$

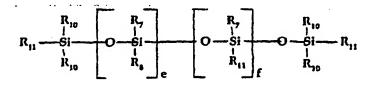
wherein:

 R_{13} , R_{14} and R_{15} are the same or different and are alkyl, substituted alkyl, alkyl aryl or alkenyl groups of up to 16 carbon atoms with the proviso that the total carbon atoms in $R_{13} + R_{14} + R_{15}$ is between

10 and 24;

with the proviso that wherein R is a mixture of organosilicone amine and organic amine moieties, at least 5 equivalent weight percent to about 60 equivalent weight percent of the total equivalent weight of amine moieties of the phospholipid composition is a quaternized organosilicone amidoamine moiety, a quaternized organosilicone tertiary amine moiety or mixtures of the same.

[0007] The silicone backbone chain, R_1 , to which the amidoamine and/or amine functional groups as herein described are attached and which are shown herein as R_{11} , corresponds to the general formula:



wherein:

R₇ and R₈, which may be the same or different are selected from alkyl, aryl, capped or uncapped polyoxyalkylene, alkaryl, aralkylene and alkenyl (vinyl);

R₁₀ can be the same or different and can be selected from alkyl, aryl and olefinic (vinyl);

which can be the same or different, can be selected from R_{10} , $-(CH_2)_{n1}$ - B_{n2} - $(CH_2)_{n3}$ -CO- NR_2 - $(CH_2)_n$ - NR_3R_4 , $-R_6$ - NR_3R_4 - and mixtures thereof, wherein R_2 , R_3 , R_4 , R_6 , R_6 , R_7 , R_8 , $R_$

above; with the proviso that at least one of R11 is an amidoamine or tertiary amine;

e can be an integer of O to 50,000;

f can be an integer of 0 to 100

10 [0008] According to a preferred embodiment, the terminal groups R₁₁ are R₁₀ and f is greater than 0.

[0009] It is evident from the above general formula for phospholipid compositions of the invention that the functional phosphorus containing group(s) can be linked terminally, laterally or both terminally and laterally to the siloxane backbone chain.

[0010] In another aspect of the present invention there is provided a method of preparing novel phospholipid compositions that may be represented by the general formula:

$$R-Y-O-P-O-P-O-A$$

wherein:

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A is selected from H, M, and R-Y-;

A₁ is selected from H, OH, OM, and R-Y-O-;

M is a cation, preferably an alkali metal;

x is O or an integer from 1 to 5;

Y is alkylene or substituted alkylene; and

R is a mixture of quaternized amidoamine and/or tertiary amine moieties as hereinabove defined;

which comprises reacting the combination of an organic amidoamine and/or organic tertiary amine reactant and a silicone modified amidoamine and/or silicone modified tertiary amine reactant with a polyphosphate, phosphite or phosphate ester reactant in the equivalent weight ratios of from about 0.7 to 3.3, of total amidoamine and/or tertiary amine reactants to 1 of polyphosphate, phosphite or phosphate ester halide reactant until the amine reactant is completely reacted, with the proviso that at least 5 equivalent weight percent to about 60 equivalent weight percent of the total equivalent weight of amine reactants will be silicone containing, said polyphosphate, phosphite or phosphate ester halide reactant being of the general formula:

wherein:

A2, is selected from H, M and X-Y-;

A, is selected from H, OH, OM, and R-Y-O-;

x is O or an integer from 1 to 5;

M is a cation, preferably alkali metal;

Y is alkylene or substituted alkylene; and

X is halogen.

[0011] The novel phospholipid compositions of the present invention comprise a class of compositions which may be represented by the general formula:

wherein:

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A is selected from H, M, and R-Y-;

A, is selected from H, OH, OM and R-Y-O-;

x is O or an integer from 1 to 5;

M is a cation, preferably an alkali metal;

may be alkylene, optionally interrupted by up to 3 oxygen atoms, of up to 12 carbon atoms, which alkylene chain may optionally be substituted with lower alkyl, alkoxyxalkyl or hydroxyalkyl, e.g., of not more than 10 carbon atoms each;

R is selected from:

a) a quaternized organosilicone amidoamine moiety of the formula:

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wherein:

R₁ is a silicone backbone chain hereinafter described to which amido amine functional group(s) as

herein described can be attached;

R₂ is hydrogen or alkyl, hydroxyalkyl or alkenyl of up to 6 carbon atoms each, or cycloalkyl of up to 6

carbon atoms, or polyoxyalkylene of up to 10 carbon atoms, preferably from 2 to 5 carbon atoms,

within the oxyalkylene unit;

R₃ and R₄, which may be the same or different, are selected from alkyl, hydroxyalkyl, carboxyalkyl of up to 6

carbon atoms in each alkyl, and polyoxyalkylene of up to 10 carbon atoms;

X- is an anion, preferably a halogen;

n is an integer from 2 to 12;

n¹ is zero or an integer from 1 to 12;

n² is 0 or 1;

n³ is an integer from 1 to 5;

is sulfur (S) or oxygen (O); with the proviso that when n² is 0, n¹ or n³ is at least 1 and when n² is

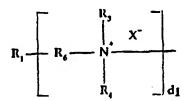
1, n1 and n3 each is at least 1; and

d is one or greater, preferably 2-10;

b) a quaternized organosilicone tertiary amine moiety of the formula:

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wherein:

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R₁ is a silicone backbone chain as hereinafter described to which amine functional groups can be at-

tached;

R₆ is alkylene, hydroxyalkylene, arylene, alkarylene, aralkylene, heteroalkylene wherein the heteroatom

can be N. S or 0 and there can be more than one of such hetero atoms in the chain;

X- is an anion, preferably a halogen;

is an integer of one or greater, preferably from 2 to 10; and

R₃ and R₄ are as hereinabove defined;

or a mixture of the quaternized organosilicone amidoamine moiety "a" and/or quaternized organosilicone tertiary amine moiety "b" above and moieties selected from the group consisting of:

c) a quaternized organic amidoamine moiety of the formula:

$$\begin{bmatrix}
O & \begin{bmatrix}
R_2 & O \\
\end{bmatrix} & \begin{bmatrix}
R_2 & C \\
\end{bmatrix} & \begin{bmatrix}
R_3 & C \\
\end{bmatrix} & \begin{bmatrix}
R_2 & R_3 & X
\end{bmatrix} & \begin{bmatrix}
R_3 & X
\end{bmatrix} & \begin{bmatrix}
R_4 & X
\end{bmatrix}$$

wherein:

R₅ is alkyl, alkenyl, alkoxyalkyl or hydroxyalkyl of from 5 to 21 carbon atoms each, alkaryl or aryl of up to 20 carbon atoms;

R₂ is hydrogen or alkyl, hydroxyalkyl or alkenyl of up to 6 carbon atoms each, cycoalkyl of up to 6 carbon atoms, or polyoxyalkylene of up to 10 carbon atoms, preferably of from 2 to 5 carbon atoms within

the oxyalkylene unit;

R₃ and R₄ which may be the same or different, are selected from alkyl, hydroxyalkyl, carboxyalkyl of up to 6 carbon atoms in each alkyl moiety, and polyoxyalkylene of up to 10 carbon atoms; in addition R, and

R₄ taken together with the nitrogen to which they are attached may represent an N-heterocycle;

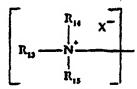
X- is an anion, preferably a halogen;

g is 0 or 1

n is integer from 2 to 12; and

n⁴ is an 1 or greater; and/or

d) an organic quaternized tertiary amine moiety of the formula:



50 wherein:

R₁₃, R₁₄ and R₁₅ are the same or different and are alkyl, substituted alkyl, alkaryl or alkenyl groups of up to

16 carbon atoms with the proviso that the total carbon atoms in $R_{13} + R_{14} + R_{15}$ is between 10 and 24;

x- is an anion, preferably a halogen.

with the proviso that wherein R is a mixture of organosilicone amine and organic amine moieties, at least 5 equivalent weight percent to about 60 equivalent weight percent of the total equivalent weight of amine moieties of the phospholipid

composition is a quaternized organosilicone amidoamine moiety, a quaternized organosilicone tertiary amine moiety or mixtures of the same.

[0012] A preferred phopholipid composition of the invention wherein Y is 2-hydroxypropylene comprises a class of compositions which may be represented by the general formula:

wherein

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A and R are as defined hereinabove.

[0013] The silicone backbone chain R_1 to which the amine functional groups as hereinabove shown are attached and which are shown herein as R_{11} , corresponds to the general formula:

$$R_{11} - Si - C - S$$

wherein:

R₇ and R₈, which may be the same or different are selected from alkyl, aryl, capped or uncapped polyoxyalkylene,

alkaryl, aralkylene and alkenyl (vinyl);

R₁₀ can be the same or different and can be selected from alkyl, aryl and olefinic (vinyl);

 R_{11} , which can be the same or different, may be selected from R_{10} , $-(CH_2)_{n1}$ - B_{n2} - $-(CH_2)_{n3}$ CO- NR_2 - $-(CH_2)_{n3}$

NR₃R₄-, -R₆-NR₃R₄- and mixtures thereof wherein R₂, R₃,R₄, R₆, B, n, n¹, n² and n³ are as defined

above; with the proviso that at least one of R₁₁ is an amidoamine or tertiary amine;

e can be an integer of 0 to 50,000;

f can be an integer of 0 to 100.

[0014] It is evident from the general formula of the novel phospholipid compositions of the invention that the functional phosphorus containing group(s) can be linked terminally, laterally or both terminally and laterally to the siloxane chain. [0015] The phospholipid compositions of the invention can be prepared by reacting corresponding silicone-modified tertiary amine and/or amidoamine reactants or combinations of corresponding silicone modified tertiary amine and/or amidoamine reactants and organic tertiary amine and/or amidoamines reactants with polyphosphate, phosphite, or phosphate ester halide reactants in appropriate stoichiometric quantities as will be described in detail hereinafter to obtain the desired products of the formula:

$$R-Y-O-P-O-P-O-A$$

and preferably

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wherein:

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- 10 A is as defined hereinabove;
 - A₁ is as defined hereinabove;
 - x is as defined hereinabove;
 - M is as defined hereinabove;
- R is selected from a quaternized organosilicone amidoamine moiety, a quaternized organosilicone tertiary amine moiety, or a mixture of moieties selected from a quaternized organosilicone amidoamine moiety, a quaternized organosilicone tertiary amine moiety, an organic amidoamine or organic tertiary amine moiety or mixtures thereof as defined hereinabove; with the proviso that wherein R is a mixture of organosilicone amine and organoamine moieties at least 5 to about 60 equivalent weight percent of the total equivalent weight of amine moieties of the phospholipid composition is a quaternized organosilicone amidoamine, a quaternized organosilicone tertiary amine and mixtures of the same.

[0016] The intermediate reactants required in the processes for preparing the phospholipid compositions of the invention can be prepared as follows:

[0017] Phosphate, polyphosphate and/or phosphite ester halide intermediate reactants based on epichlorhydrin can be prepared by known procedures illustrated as follows: I

wherein:

A2 is selected from H, M and X-Y-;

A₃ is selected from H, OH, OM and X-Y-O-;

A4 is H, OM or OH;

A₅ is H or M;

a is from 0.5 to 3.5, preferably 1 to 3;

b is from 1 to 3, preferably 1-2;

X is 0 or an integer from 1-5;

M is a cation, preferably alkali metal;

x is halogen;

Y is 2 hydroxyproplyene.

[0018] The above coupling reaction is carried out in an aqueous media, preferably in the range of 30-50% concentration, having a pH range of 5.0-8.0.

II

a CII_CII_X +
$$110 - P - 0$$
 $P - 0$ $P - 0$

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$$X-Y-O-P-O-P-O-A_{\delta}$$

wherein:

20 A₇ is H, OH or X-Y-O-;

A_R is H or -Y-X;

a is from about 0.5 to 7, preferably, from about 1 to 3;

is 0 or an integer from 1-5;

X is halogen;

25 Y is 2 hydroxypropylene.

[0019] The reaction is preferably carried out in absence of water with slight excess epichlorohydrin.

[0020] Phosphate, phosphite and/or polypliosphate ester intermediate reactants for preparing phosphobetaine, pyrophosphobetaine and the like compositions of the invention can also be prepared by known procedures such as are disclosed, for example, in U.S. -A- 4,617,414.

[0021] Silicone-modified amidoamine intermediate reactants suitable for use in preparing the phospholipid composition of the invention can be prepared as follows:

[0022] Also suitable as phosphate and phosphite intermediate reactants are such reactants prepared by known procedures illustrated as follows:

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(3-a)
$$X-R_{16}$$
 OH + POCl₃ \longrightarrow ($X-R_{16}$ O $\frac{1}{3-a}$ P-Cl_a + (3-a) HC

45 III (b)

$$(X-R_{10} - O) = P - CI_{0} - P - CI_{0} -$$

50 wherein

a is O or an integer from 1 to 2; X is halogen, preferably bromine; R₁₆ is alkylene.

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IV (a)
$$\begin{array}{c}
CI \\
X - R_{16} - OH + PCI_{3} \longrightarrow X - R_{16} - O - P - CI + HC
\end{array}$$

wherein

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X is a halogen, preferably bromine; R₁₆ is alkylene.

[0023] Carrying out reactions III(a) and IV(b) in presence of a tertiary amine HCI acceptor is preferred to prevent formation of free acid.

[0024] Silicone-modified amidoamine intermediate reactants suitable for use in preparing the phospholipid composition of the invention can be prepared as follows:

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$$R_{1} = \left(\frac{CH_{2}}{n_{1}} \frac{B_{1}}{n_{2}} + \frac{CH_{2}}{n_{3}} \frac{C}{C} - OR_{12} \right) + D = R_{2} + R_{3}$$

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$$R_{1} = \left(\frac{CH_{2}}{n_{1}} \frac{B_{1}}{n_{2}} + \frac{CH_{2}}{n_{3}} \frac{C}{n_{3}} - OR_{12} \right) + D = R_{2} + CH_{2} +$$

wherein:

R₁ is a silicone backbone chain as herein defined;

R₂ is as hereinabove defined; R₃ and R₄ is as previously defined;

 R_3 and R_4 is as previously defined is hydrogen or alkyl;

B is sulfur or oxygen; with the proviso that when n^2 is 0, n^1 or n^3 is at least 1 and when n^2 is 1, n^1 and n^3

each is at least 1;

n is an integer from 2 to 12;

55 n¹ is zero or an integer of one or greater;

n² is 0 or 1;

n³ is an integer of 1 to 5;

d and D is an integer from 1 or greater, generally from 1-50 and preferably 2-10. The reactant ratio of the amine

reactant to the carboxyl reactant on the silicon is preferably 1:1 but can be varied in ratio of 0.8 - 1.2.

[0025] The above coupling reaction (V) for preparing the silicone-modified amidoamine intermediate reactants can be carried out neat or can be carried out in an inert solvent such as xylene, toluene, benzene, chlorobenzene or the like.

[0026] The polysiloxane-containing functional carboxylic acids or derivatives thereof (terminal, lateral or combination of terminal and lateral) applicable for use in preparing the silicone-modified amidoamine intermediate reactants as set forth in the reaction sequence illustrated above (III) can be prepared by a variety of known procedures such as illustrated by the following:

VI

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wherein:

h is an integer from 1-100; j is an integer from 0-1000.

55 wherein:

k is an integer from 1-1000.

VIII

wherein:

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I is an integer from 1-100; m is an integer from 0-1000.

[0027] Suitable carboxyl functional silicone compositions having terminal, lateral or combinations of terminal and lateral functional groups are available commercially, for example, from Shin-Etsu. While the molecular weight of the silicone compositions which may be employed are not critical, and suitable compositions may have amine equivalent weights of 8000, or even higher, silicone compositions having amine equivalent weights from about 1500 to about 6000 are in general preferred.

[0028] Silicone-modified tertiary amine intermediate reactants which are suitable for use in preparing alternate embodiments of the silicone-modified phospholipid compositions of the invention can be silicone-modified tertiary amines (terminal, lateral or combinations thereof) which are prepared by a variety of known procedures such as disclosed, for example, in U.S. -A- 3,389,160 which describes the preparation of a carbinol containing tertiary silicone amine encompassing the reaction of a secondary amine with an epoxy containing silicone fluid (example 1) and by Snow et al, J. Langmuir, 1990, 6(2), pp 336-39, wherein the preparation of tertiary amino functional siloxanes result from the hydrosilylation of olefinic tertiary amines with hydride siloxane fluids employing a platinum catalyst.

[0029] In addition, the preparation of a suitable functional tertiary amino alkyl dimethylsilyl capped material is disclosed in U.S.-A-4,918,210, at example 1, part 2 which consists of the Pt catalyzed addition of a terminal hydride containing silicone fluid with N-allyl-diethylamine. In general, silicone containing tertiary amine intermediate reactants with molecular weights ranging, between about 1000 and 6000 are most advantageously employed.

[0030] The organic amidoamine intermediate reactants suitable for use in preparing the phospholipid compositions of the invention can be prepared as follows:

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$$R_{s} = C = \begin{bmatrix} R_{2} & O \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{12} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{12} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2} \\ -N & -C \end{bmatrix}_{n_{d}} = C = \begin{bmatrix} R_{2} & R_{2}$$

wherein:

 R_5 is as hereinabove defined; R_2 is as hereinabove defined; R_3 and R_4 is as hereinabove defined; R_{12} is hydrogen or alkyl; g is 0 or 1; n is an integer from 2 to 12; and n^4 is 1 or greater.

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[0031] The organic amidoamine intermediate reactants suitable for use in preparing the phospholipid compositions shown in the above coupling reaction are known or are generally prepared in accordance with conventional techniques. A wide variety of commercially available tertiary amino alkyl amines are suitable for use in reaction with an acid or acid derivative to prepare suitable amidoamines, as are the amidoamines themselves. The suitable tertiary amino alkyl amines can be primary or secondary amines with the proviso that the total number of carbons in the acid portion of the molecule be greater than 6, i.e. to give a hydrophobic moiety necessary for surface activity properties. Suitable amidoamines include acyl derivatives of aminoacid products such as glycine and sarcosine (N-methylglycine) including for example, products available under the Tradename HAMPOSYL from the Hampshire Chemical Co.

[0032] The organic tertiary amine reactants suitable for use in preparing the phospholipid compositions of the invention can be prepared using procedures well known in the art and many suitable compositions are available.

[0033] Exemplary tertiary amines include:

tributylamine
bis(hydroxyethyl)hexylamine
bis(2-hydroxyethyl)cocoamine
N,N-dimethyl-dodecylamine
N,N-dimethyl-tetradecylamine
N,N-dimethyl-hexadecylamine
N,N-dimethyl-cocoamine

N,N-dimethyl-cetylamine dimethyl (C₈-C₁₆) alkyl amine. N,N-dimethyl-octadecylamine

[0034] As indicated, the phospholipid compositions of the invention can be prepared by reacting a tertiary amine or amidoamine functional silicones or combination of tertiary amine and/or amidoamine functional silicones and organic tertiary amine and/or amidoamine reactants with the phosphate ester halide reactants herein described in appropriate stoichiometric quantities as will be discussed in detail hereinafter.

[0035] The reaction of silicone containing amidoamine or tertiary amine functional groups with phosphate, polyphos-

phate and/or phosphite ester halide reactants in molar equivalents from about 0.7 to 3.3 of the amine functional silicones to 1 of the phosphate, phosphite and/or polyphosphate ester halide reactants based on the reaction can be readily carried out in an aqueous or aqueous organic co-solvent reaction systems wherein the number of grams of silicone fluid containing an amine equivalent lie in the area up to about 1200. Reactions will go to completion as demonstrated by chlorine analysis, alkali number titration, and homogeneity of reaction. When the number of grams of fluid per amine equivalent weight of the silicone reactants is greater than about 1200 to 2000 the reactants are partially or completely insoluble in the reaction system and an incomplete reaction will result. With silicone composition reactants having amine equivalent weights above about 2000, the addition of a co-solvent to an aqueous reaction system will not increase the solubility of the reactants as evidenced by phase separation and/or other signs of incomplete reaction. Surprisingly and unexpectedly, it has been found that phospholipid compositions of the invention can be prepared in substantial complete reaction form and in completely soluble reaction systems using tertiary amine and/or amidoamine functional silicone reactants having amine equivalent weights of 6000, or even greater, by also incorporating in the reaction system, organic tertiary amine and/or amidoamine reactants as herein described in conjunction with the functional silicone reactants. The organic tertiary amine and/or amidoamine reactant is added to the reaction system as a partial replacement of a molar equivalent amount of the functional silicone reactant, which substantially maintains the above noted molar equivalent ratios of amine reactants to phosphate ester halide reactants in the reaction mixture.

[0036] The reaction of a combination of organic tertiary amine and/or amidoamine reactants and silicone modified amine reactants with phosphate, phosphite and/or polyphosphate ester halide reactants will proceed to completion at an elevated temperature, preferably a temperature ranging from about 75°C to 95°C, with the formation of the silicone-modified phospholipid compositions. The order of addition of the reactants is not critical and while a heterogeneous mixture may result when all reactants are admixed, the system becomes homogenous as the reaction proceeds. The reaction may start slowly while the mixture is heterogeneous but the reaction mixture will become substantially clear as the reaction proceeds. In accordance with the process of the invention, silicone-modified phospholipid compositions which contain at least 5 equivalent weight percent to about 60 equivalent weight percent silicone of the total amine containing moieties can be prepared, which silicone-containing phospholipid compositions will be completely soluble in aqueous/solvent or, preferably, aqueous systems while exhibiting surface active properties including low surface tension, high foaming and substantivity characteristics, low ocular and skin irritation and the like. Thus, it is possible by choice of particular amine functional silicone, organic amine and phosphate ester halide reactants to obtain soluble and preferably aqueous soluble silicone-modified phospholipid composition with a wide range of surface active agent properties for use in a variety of applications.

[0037] The preparation of specific composition of the invention is illustrated by the following specific examples which are provided herein for purposes of illustration only and are not intended to limit the scope therein.

EXAMPLE 1

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[0038] A carboxyl containing (pendant), trimethylsilyl capped silicone fluid obtained from Shin-Etsu under the designation X-22310 having a carboxyl content of 3.5% and an initial equivalent weight of 1289 is used in this example. [0039] 1592.7 grams (1.235 moles) of the above polysiloxane fluid is combined with 189 grams (50% excess) of dimethylaminopropyl amine (DMAPA) in a reaction vessel and heated slowly under nitrogen to about 185°C. The reaction is carried out for 4 hours while volatiles come off and are collected. There is collected a total of 44ml of volatiles as two layers, the bottom layer of 25ml is water with some volatilized dimethylaminopropyl amine. The residue in the reaction vessel has an acid number less than 1.

[0040] The materials are then heated to a temperature of 120°-140°C under a vacuum of 1330 Pa (10mm) for 3 hours. At this point the polysiloxane/DMAPA reaction product is found to have an alkali number of 42 and equivalent weight of 1335.

[0041] 264 grams (0.9428 moles) of cocoyl sarcosine obtained under the tradename HAMPOSYL C from Hampshire Chemical Corp. is reacted with 144 grams (an excess) of dimethylaminopropyl amine in 150ml of refluxing xylene under a nitrogen atmosphere, with water being removed as it is formed. After 4 hours, water no longer evolves and the xylene and any volatiles are removed by heating the reaction mixture to 130°C at 1330 Pa (10mm) vacuum for 3 hours. 348 grams of a product having an alkali number of 185 is obtained.

[0042] 2.7 grams (0.002 mole) of the above silicone/DMAPA product are combined with 2.4 grams (0.008 moles) of the above sarcosine/DMAPA product; 3.12 grams (0.0033 moles) of a 40% concentration of a phosphate ester halide reactant prepared by the reaction of 3 moles of epichlorohydrin with one mole of 85% phosphoric acid in the presence of one mole sodium hydroxide; and 13 grams water. The amidoamine reactants are used in an equivalent weight ratio of 4:1 of organic amidoamine to silicone amidoamine reactants.

[0043] The reaction mixture is heated for 3 hours at 90°C at which time a hazy but homogeneous reaction product is obtained having a % NaCl content of 2.6 (2.8 theoretical).

[0044] The resultant product when added to water forms a clear solution which foamed well, as compared to the

silicone fluid starting material which forms a hazy, non-foaming mixture in water.

EXAMPLE 2

[0045] 6.67 grams (0.005) of a silicone/DMAPA reaction product (equivalent weight 1335) prepared as described in example 1 is mixed with 3.84 grams (0.010 mole) of the reaction product of linoleic acid and dimethylaminopropyl amine prepared by well known, conventional techniques, 4.69 grams (0.005 mole) of phosphate ester halide reactant prepared as described in example 1, and 26 grams of water. The reaction mixture is heated at 90°C for 3 hours at which time a clear, viscous, yellow solution is obtained having a % NaCl content of 1.9% (2.1 theoretical). 10

[0046] The resulting product added to water forms a clear solution which produces a stable foam.

EXAMPLE 3

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[0047] A carboxyl containing (pendant) trimethylsilyl capped silicone fluid obtained from Shin Etsu under the designation X-223701E having a carboxyl content of 0.95% and initial equivalent weight of 4740 is used in this example. [0048] 967 grams (0.204 mole) of the above silicone fluid is combined with 31.2 grams (excess) of dimethylaminopropyl amine (DMAPA) and heated slowly under nitrogen to about 185°C while volatiles coming off are collected. After a period of 4 hours, 15.5ml of volatiles are collected as two layers, the bottom one being water with some volatilized DMAPA.

[0049] The reaction mixture having an acid number less than 1 is then heated to about 140°C at 665 Pa (5mm) vacuum for 4 hours to produce a product having an alkali number of 11.4 and an equivalent weight of 4921. [0050] 9.8 grams (0.002 mole.) of the above silicone/DMAPA reaction product are mixed with 2.4 grams (0.008 mole) of the cocyl sarcosine/DMAPA reaction product prepared as in example 1, 3.13 grams (0.0033 mole) of the phosphate ester halide reactant of example 2 and 29.5 grams water (30% solids concentration). The reaction mixture is heated for 3 hours at 94° to 95°C during which time a clear but somewhat hazy solution is obtained. The % NaCl content of the reaction product is 1.4. The resulting product forms a clear solution when added to water which produces a large amount of stable foam.

EXAMPLE 4

[0051] Using the silicone/DMAPA reaction product of example 3 (equivalent weight of 4921) and the phosphate ester halide reactant of example 1, a series of mixtures of the two reactants in equivalent weight ratios ranging from 0.1 to 6:1 in water at solid content concentrations ranging from 10-70% are reacted. The various combination of reactants failed to form a homogeneous reaction mixture during reaction times ranging from 2 to 10 hours at temperatures up to 100°C. Similar results are obtained with the addition of cosolvents such as isopropanol and propylene glycol to the reaction system.

EXAMPLE 5

[0052] A reaction mixture is prepared from 39.2 grams of the silicone/DMAPA reaction product of example 3, 12 40 grams of a linoleic acid/DMAPA reaction product, 15.92 grams of the cocoyl sarcosine/DMAPA reaction product of example 1, 28.2 grams of the phosphate ester halide reactant of example 1 and 91.6 grams of water. The reaction mixture is heated to 90°C for four hours during which time a clear solution is formed having a % NaCl content of 2.7.

EXAMPLE 6

[0053] Trisiloxane 1,1,1,3,5,5,5-hepta methyl, 3-Dimethylamino propyl trisiloxane prepared by the platinum catalyzed addition of the corresponding heptamethyltrisiloxane to N,N Dimethyl allyl amine as described in U.S. -A- 3,658,867 is used in this example. A reaction mixture containing 2.4 grams of the above trisiloxane (0.008 mole), 3.2 grams (0.012 mole) of cetyldimethylamine, 6.24 grams of the phosphate ester halide reactant of example 1, and 15 ml water is prepared and heated to 90°C for 3 hours. A homogeneous, clear solution is formed having an alkali number less than 2. [0054] A few drops of the product when added to 100 ml of water forms a clear solution which forms a stable foam when shaken. Although some preferred embodiments have been described, many modifications and variations may be made thereto in light of the above teachings.

EXAMPLE 7

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[0055] DiSodium 1,3 Bis 3 chloro-2 hydroxy propyl pyrophosphate is prepared by charging 446 Parts of Na₄P₂O₇

10H₂O (1 mole) and 17B parts H₄P₂O₇ (1 mole) with 1000 parts deionized water to a reaction vessel and reacting the same with 320 parts epichlorohydrin at 60-80°C for 3-4 hours. 81.4 parts (0.2 equivalents) of the above reaction product is combined with a mixture of a pendant trimethylsilyl silicone amidoamine having an equivalent weight of 1335 (0.05 equivalent) and 57.6 parts of N-Dimethylaminopropyl linoleamide (0.15 equivalents) and then diluted with 480 grams of water to a 30% concentration. After the solution is adjusted to a pH of 8, the reaction mixture is heated to 90-95°C for a period of 4-5 hours at which time a clear solution forms. The reaction is monitored via argentometric estimation for covalent chloride to ionic chloride and the reaction is completed in 5 hours.

EXAMPLE 8

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[0057] A 3-chloro-2 hydroxy propylester salt of phosphorous acid is prepared by charging 41 grams (0.5 mole) of phosphorus and, 409 grams of water and 50 grams of a 50% NaOH solution (0.6 mole) to a reaction vessel and warmed to 75°C. 46.25 grams (0.5 mole) of epichlorohydrin is then added and the reaction mixture is heated at 75°C for 1 ½ hours with stirring.

[0058] 196 parts of the combined epichlorohydrin-phosphite reaction mixture is admixed with a 50% sodium hydroxide solution to achieve a pH of 8 followed by adding the combination a pendant trimethylsilyl capped silicone amidoamine having an equivalent weight of 1335 (13.35 gram) and 27.3 parts of cocoyl sarcosine amidoamine (0.09 equivalents). The reaction mixture is diluted with water to achieve 25% solids.

[0059] The reaction mixture is heated to 90°C for 2 hours whereupon a clear aqueous solution is formed.

EXAMPLE 9

[0060] 3-Bromopropyl diacid phosphate (BrCH₂CH₂OPO(OH)₂) is prepared by reacting 3-Bromopropanol with POCl₃ utilizing one equivalent of triethyamine in methylene chloride solvent. The Dichloride is isolated and hydroyzed to the diacid with water. The pH of the product in water is adjusted to 8 followed by the addition of an equivalent amount of the 50:50 combination of a trimethylsilyl capped silicone amidoamine having an equivalent weight of 1335 and N-Dimethyl aminopropyl derivative of linoleamide. The reaction mixture is adjusted with water to 30% total solids.

[0061] The reaction mixture is heated for 4 hours at 90°C. A clear solution is formed.

Claims

1. Phospholipid compositions of the formula

$\begin{array}{c|c} \mathbf{R} - \mathbf{Y} - \mathbf{O} & \mathbf{O} & \mathbf{O} \\ \mathbf{A}_{1} & \mathbf{O} & \mathbf{A}_{2} \end{array}$

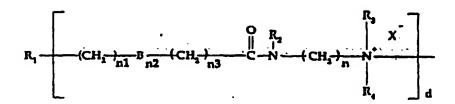
characterized in that

A is selected from H, M and R-Y-; A₁ is selected from H, OH, OM and R-Y-O-; x is 0 or an integer from 1 to 5; M is a cation;

Y is alkylene or substituted alkylene;

R is a quaternized organosilicone amine moiety selected from the group consisting of:

a) a quaternized organosilicone amidoamine moiety of the formula:



wherein:

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R₁ is a silicone backbone chain to which amidoamine functional group(s) can be attached and is represented by the formula:

 $R_{j_{1}} = \begin{bmatrix} R_{j_{0}} & R_{j_{0}} & R_{j_{0}} \\ \vdots & \vdots & \vdots \\ R_{j_{0}} & R_{j_{0}} \end{bmatrix} = \begin{bmatrix} R_{j_{0}} & R_{j_{0}} \\ \vdots & \vdots & \vdots \\ R_{j_{1}} & R_{j_{0}} \end{bmatrix} = \begin{bmatrix} R_{j_{0}} & R_{j_{0}} \\ \vdots & \vdots & \vdots \\ R_{j_{1}} & R_{j_{0}} \end{bmatrix}$

wherein:

R₇ and R₈, which may be the same or different are selected from alkyl, aryl, capped or uncapped polyoxy-

alkylene, alkaryl, aralkylene or alkenyl (vinyl);

R₁₀ can be the same or different and is selected from alkyl, aryl or olefinic (vinyl);

 R_{11} , which can be the same or different, is selected from R_{10} , $-(CH_2)_{n1}$ - B_{n2} - $(CH_2)_{n3}$ - $CO-NR_2$ - $(CH_2)_n$ -

 NR_3R_4 , $-R_6-NR_3R_4$ - or mixtures thereof, wherein R_2 , R_3 , R_4 , R_6 ,

defined above; with the proviso that at least one of \mathbf{R}_{11} is an amidoamine;

e can be an integer of 0 to 50,000;

f can be an integer of 0 to 100.

R₂ is hydrogen or alkyl, hydroxyalkyl or alkenyl of up to 6 carbon atoms each, or cycloalkyl of up

to 6 carbon atoms, or polyoxyalkylene of up to 10 carbon atoms within the oxyalkylene unit;

R₃ and R₄, which may be the same or different, are selected from alkyl, hydroxyalkyl, carboxyalkyl of up

to 6 carbon atoms in each alkyl, or polyoxyalkylene of up to 10 carbon atoms;

X- is an anion, preferably a halogen;

n is an integer from 2 to 12;

n¹ is zero or an integer from 1 to 12;

n² is 0 or 1;

n³ is an integer from 1 to 5;

B is sulfur (S) or oxygen (O); with the proviso that when n' is 0, n¹ or n³ is at least 1 and when n²

is 1, n1 and n3 each is at least 1; and

d is one or greater;

 \cdot b) a quaternized organosilicone tertiary amine moiety of the formula:

 $R_{1} = \begin{bmatrix} R_{0} & X \\ & N & X \end{bmatrix}_{d_{1}}$

wherein:

R₁ as herein above defined is a silicone backbone chain to which amidoamine or amine functional groups can be attached;

R₆ is alkylene, hydroxyalkylene, arylene, alkarylene, aralkylene, heteroalkylene wherein the heteroatom can be N, S or O and there can be more than one of such hetero atoms in the chain;

X- is an anion;

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d¹ is an integer of one or greater;

R₃ and R₄ are as hereinabove defined; and mixtures of the same.

- 2. The phospholipid composition according to claim 1, characterized in that f is 0.
- 3. The phospholipid composition according to claim 1 characterized in that the terminal groups R₁₁ are R₁₀ and f is greater than 0.
- 4. The phospholipid composition according to claim 1 characterized in that Y is -CH₂-CHOH-CH₂-.
- 5. Phospholipid compositions represented by the following general formula:

characterized in that

A is selected from H, M, or R-Y-;

A, is selected from H, OH, OM or R-Y-O-;

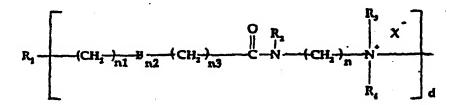
x is 0 or an integer from 1 to 5;

M is a cation:

Y is alkylene or substitued alkylene; and

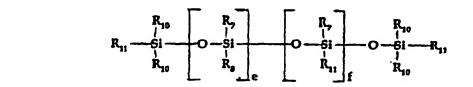
R is a mixture of quaternized organosilicone amine moieties selected from the group consisting of:

a) a quaternized organosilicone amidoamine moiety of the formulas:



wherein:

is a silicone backbone chain to which amidoamine or amine functional group(s) can be attached and is represented by the formula:



wherein:

R₇ and R₈, which may be the same or different are selected from alkyl, aryl, capped or un-

capped polyoxyalkylene, alkaryl, aralkylene or alkenyl (vinyl);

 R_{10} can be the same or different and is selected from alkyl, aryl or olefinic (vinyl); R_{11} , which can be the same or different, is selected from R_{10} , -(CH₂)_{n1}- R_{n2} -(CH₂)_{n3}- R_{n2} -(CH₂)_{n3}

CO-NR₂-(CH₂)_n-NR₃R₄, -R₆-NR₃R₄- or mixtures thereof, wherein R₂, R₃, R₄, R₆, B, n, n¹, n² and n³ are as defined above; with the proviso that at least one

of R₁₁ is an amidoamine;

e can be an integer of 0 to 50,000;

f can be an integer of 0 to 100;

R₂ is hydrogen or alkyl, hydroxyalkyl or alkenyl of up to 6 carbon atoms each, or cycloalkyl of up to 6 carbon atoms, or polyoxyalkylene of up to 10 carbon atoms within the oxyalkylene unit;

R₃ and R₄, which may be the same or different, are selected from alkyl, hydroxyalkyl, carboxyalkyl of up to 6 carbon atoms in each alkyl, or polyoxyalkylene of up to 10 carbon atoms; in addition R₃ and R₄ taken together with the nitrogen to which they are attached represent a N-heterocycle;

X is an anion:

n is an integer from 2 to 12;

n¹ is zero or an integer from 1 to 12;

n² is 0 or 1:

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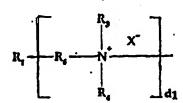
n³ is an integer from 1 to 5;

B is sulfur (S) or oxygen (O); with the proviso that when n² is 0, n¹ or n³ is at least 1 and when

n2 is 1, n1 and n3 each is at least 1; and

d is one or greater; and

b) a quaternized organosilicone tertiary amine moiety of the formula:



wherein:

R₁ as herein above defined is a silicone backbone chain to which amidoamine or amine functional

groups can be attached;

R₆ is alkylene, hydroxyalkylene, arylene, alkarylene, aralkylene, heteroalkylene wherein the heteroatom can be N, S or O and there can be more than one of such hetero atoms in the chain;

X- is an anion;

d¹ is an integer of one or greater;

R₃ and R₄, which may be the same or different, are selected from alkyle, hydroxyalkyl, carboxyalkyl of up to 6 carbon atoms in each alkyl, or polyoxyalkylene of up to 10 carbon atoms; in addition R,

and R4 taken together with the nitrogen to which they are attached represent a N-heterocycle;

and a quaternized organic amine moiety selected from the group consisting of:

c) a quaternized organic amidoamine moiety of the formula:

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$$\begin{bmatrix} O & \begin{bmatrix} R_2 & O \\ & & \end{bmatrix} & \begin{bmatrix} R_2 & R_3 & X^- \\ & & \end{bmatrix} \\ R_3 - C & \begin{bmatrix} CH_1 \\ N \end{bmatrix} & \begin{bmatrix} CH_2 \\ N \end{bmatrix} & \begin{bmatrix} CH_2 \\ N \end{bmatrix} & \begin{bmatrix} R_4 \\ & \end{bmatrix} \end{bmatrix}$$

10 wherein:

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R₅ is alkyl, alkenyl, alkoxyalkyl or hydroxyalkyl of from 5 to 21 carbon atoms each, alkaryl or aryl of

up to 20 carbon atoms;

R₂ is hydrogen or alkyl, hydroxyalkyl or alkenyl of up to 6 carbon atoms each, cycloalkyl of up to 6

carbon atoms, or polyoxyalkylene of up to 10 carbon atoms within the oxyalkylene unit;

R₃ and R₄ which may be the same or different, are selected from alkyl, hydroxyalkyl, carboxyalkyl of up to 6 carbon atoms in each alkyl moiety, and polyoxyalkylene of up to 10 carbon atoms; in addition

R₃ and R₄ taken together with the nitrogen to which they are attached represent an N-hetero-

cycle;

X is an anion;

g is 0 or 1;

n is integer from 2 to 12; and

n⁴ is 1 or greater; and

d) an organic quaternized tertiary amine moiety of the formula:

wherein:

R₁₃, R₁₄ and R₁₅ are the same or different and are alkyl, substituted alkyl, alkyl aryl or alkenyl groups of up to 16 carbon atoms with the proviso that the total carbon atoms in R₁₃ + R₁₄ + R₁₅ is between 10 and 24;

with the proviso that wherein R is a mixture of organosilicone amine and organic amine moieties, at least 5 equivalent weight percent to about 60 equivalent weight percent of the total equivalent weight of amine moieties of the phospholipid composition is a quaternized organosilicone amidoamine moiety, a quaternized organosilicone tertiary amine moiety or mixtures of the same.

- 6. The phospholipid compositions according to claim 5, characterized in that g in the quaternized organic amidoamine moiety c) is 1.
- 7. The phospholipid composition according to claim 5 characterized in that f is 0.
- 8. The phospholipid composition according to claim 5, characterized in that the terminal groups R₁₁ are R₁₀ and f is greater than 0.
- 9. The phospholipid compositions according to Claim 5, characterized in that R is a mixture of:
 - a) quaternized organosilicone amidoamine moieties of the formula:

$$R_{1} = \begin{bmatrix} O & R_{2} & R_{3} & R_{4} & R_{4}$$

10 wherein:

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R₁ as defined in claim 5 is a silicone backbone chain to which amidoamine and/or amine functional

group(s) can be attached;

R₂ is hydrogen or alkyl, hydroxyalkyl or alkenyl of up to 6 carbon atoms each, or cycloalkyl of up

to 6 carbon atoms, or polyoxyalkylene of up to 10 carbon atoms within the oxyalkylene unit;

R₃ and R₄, which may be the same or different, are selected from alkyl, hydroxyalkyl, carboxyalkyl of up

to 6 carbon atoms in each alkyl, or polyoxyalkylene of up to 10 carbon atoms;

X- is an anion;

n is an integer from 2 to 12;

n¹ is zero or an integer from 1 to 12;

n² is 0 or 1;

n³ is an integer from 1 to 5;

B is sulfur (S) or oxygen (O); with the provision that when n² is 0, n¹ or n³ is at least 1 and when

n2 is 1, n1 and n3 each is at least 1; and

d is one or greater; and

c) quaternized organic amidoamine moieties of the formula:

$$\begin{bmatrix}
O & R_2 & O \\
R_3 - C & N - (CH_3)_{n4} & C \\
\end{bmatrix} \begin{bmatrix}
R_2 & R_3 & X - \\
N - (CH_2)_{n} & N \\
g & R_4
\end{bmatrix}$$

wherein:

R₅ is alkyl, alkenyl, alkoxy, alkyl or hydroxyalkyl of from 5 to 21 carbon atoms each, alkaryl or aryl

of up to 20 carbon atoms;

R₂ is hydrogen or alkyl, hydroxyalkyl or alkenyl of up to 6 carbon atoms each, cycloalkyl of up to 6 carbon atoms, or polyoxyalkylene of up to 10 carbon atoms within the oxyalkylene unit;

R₃ and R₄ which may be the same or different, are selected from alkyl, hydroxyalkyl, carboxyalkyl of up to

6 carbon atoms in each alkyl moiety, or polyoxyalkylene of up to 10 carbon atoms; in addition

R, and R_4 taken together with the nitrogen to which they are attached represent an N-heterocycle;

X- is an anion;

g is 0 or 1;

n is integer from 2 to 12;

n⁴ is 1 or greater.

with the proviso that at least 5 equivalent weight percent to about 60 equivalent weight percent of the total equivalent weight of amine moieties of the phospholipid composition is a quaternized organosilicone amidoamine moiety or mixtures of the same.

10. A method of preparing novel phospholipid compositions according to Claim 5 represented by the general formula:

characterized in that

A is selected from H, M, or R-Y-;

A, is selected from H, OH, OM or R-Y-O-;

x is 0 or the intefer from 1 to 5;

M is a cation;

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Y is alkylene or substituted alkylene; and

R is a mixture of quaternized amidoamine and/or tertiary amine moieties as hereinabove defined;

which comprises reacting the combination of an organic amidoamine organic tertiary amine reactant or mixture of the same and a silicone modified amidoamine or silicone modified tertiary amine reactant or mixtures of the same with a polyphosphate, phosphite or phosphate ester halide reactant in the equivalent weight ratios of from about 0.7 to 3.3, of total amidoamine or tertiary amine reactants to 1 of polyphosphate, phosphite or phosphate ester halide reactant until the amine reactant is completely reacted, with the proviso that at least 5 equivalent weight percent to about 60 equivalent weight percent of the total equivalent weight of amine reactants will be silicone containing, said polyphosphate, phosphite or phosphate ester halide reactant being of the general formula:

 $X - Y - O - P - O - P - O - A_2$ $A_3 - C - O - A_3$

wherein:

 A_2 is selected from H, M and X-Y-; A_3 is selected from H, OH, OM and X-Y-O-;

x is 0 or the integer from 1 to 5;

M is a cation;

Y is alkylene or substituted alkylene; and

X is halogen.

Patentansprüche

1. Phospholipid-Zusammensetzungen gemäß der Formel

R-Y-O-P-O-A

dadurch gekennzeichnet, dass

A ausgewählt wird unter H, M und R-Y-;

A₁ ausgewählt wird unter H, OH, OM und R-Y-O-;

- x für 0 oder für eine ganze Zahl von 1 bis 5 steht;
- M ein Kation ist;
- 5 Y ein Alkylen oder ein substituiertes Alkylen ist;
 - R eine quaternierte Organosilikon-Aminhälfte darstellt, die ausgewählt wird aus der Gruppe bestehend aus:
 - a) einer quaternierten Organosilikon-Amidoaminhälfte gemäß der Formel:

in welcher:

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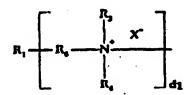
R₁ eine Silikonhauptkette ist, an welche eine oder mehrere Funktionsgruppen von Amidoamin angehängt werden können und welche wiedergegeben wird durch die Formel:

 $R_{11} = \begin{bmatrix} R_{10} & R_{7} & R_{7} & R_{10} & R_{11} & R_{10} & R_{11} & R_{10} & R_{11} & R_{10} & R_{11} & R_{10} & R$

- in welcher:
- R₇ und R₈, die identisch oder unterschiedlich sein können, ausgewählt werden unter Alkyl, Aryl, verkapptem oder unverkapptem Polyoxyalkylen, Alkaryl, Aralkylen oder Alkenyl (Vinyl);
- R₁₀, identisch oder unterschiedlich sein kann, und ausgewählt wird unter Alkyl, Aryl oder olefinisch (Vinyl) ist;
- R₁₁, das identisch oder unterschiedlich sein kann, ausgewählt wird unter R₁₀, (CH₂)_{n1}-B_{n2}-(CH₂)_{n3}-CO-NR₂-(CH₂)_n-NR₃R₄, -R₆-NR₃R₄- oder einer Mischung derselben, worin R₂, R₃, R₄, R₆, B, n, n¹, n², und n³ übereinstimmen mit dem was unten definiert ist; unter der Bedingung, dass mindestens ein R₁₁ ein Amidoamin ist;
- e eine ganze Zahl von 0 bis 50.000 sein kann;
- f eine ganze Zahl von 0 bis 100 sein kann;
- R₂ Wasserstoff oder Alkyl, Hydroxyalkyl oder Alkenyl mit bis zu 6 Kohlenstoffatomen ein jedes darstellt, oder Cycloalkyl mit bis zu 6 Kohlenstoffatomen, oder Polyoxyalkylen mit bis zu 10 Kohlenstoffatomen innerhalb der Oxyalkyleneinheit;
- R₃ und R₄, die identisch oder unterschiedlich sein können, ausgewählt werden unter Alkyl, Hydroxyalkyl, Carboxyalkyl mit bis zu 6 Kohlenstoffatomen in einem jeden Alkyl, oder Polyoxyalkylen mit bis zu 10 Kohlenstoffatomen;

	Χ-	ein Anion ist, vorzugsweise ein Halogen;
	n	für eine ganze Zahl von 2 bis 12 steht;
5	n¹	für Null oder für eine ganze Zahl von 1 bis 12 steht;
	n ²	für 0 oder für 1 steht;
10	n ³	für eine ganze Zahl von 1 bis 5 steht;
	В	Schwefel (S) oder Sauerstoff (O) darstellt; unter der Bedingung dass wenn n² gleich 0 ist, n¹ oder n³ mindestens 1 ist, und dass wenn n² gleich 1 ist, n¹ und n³ ein jedes mindestens 1 beträgt;
15	d	für eins oder für mehr steht;

b) einer quaternierten Organosilikon-tert-Aminhälfte gemäß der Formel:



in welcher

	III Welchei	
30	R ₁ ,	wie es oben definiert ist, eine Silikonhauptkette darstellt, an welche Funktionsgruppen von Amidoamin oder Amin angehängt werden können;
35	R ₆ ,	Alkylen, Hydroxyalkylen, Arylen, Alkarylen, Aralkylen, Heteroalkylen darstellt, wobei das Heteroatom N, S oder O sein kann und es kann sich mehr als ein solches Heteroatom in der Kette befinden;
	X -	ein Anion ist;
40	d ₁ .	für eine ganze Zahl von 1 oder mehr steht;
40	R ₃ und R ₄ ,	stimmen überein mit dem was oben definiert ist; ebenfalls stellen sie Mischungen derselben dar.

- Phospholipid-Zusammensetzung gemäß Anspruch 1, dadurch gekennzeichnet, dass f gleich 0 ist.
- Phospholipid-Zusammensetzung gemäß Anspruch 1, dadurch gekennzelchnet, dass die Endgruppen R₁₁ aus R₁₀ bestehen und f größer als 0 ist.
 - Phospholipid-Zusammensetzung gemäß Anspruch 1, dadurch gekennzeichnet, dass Y aus -CH₂-CHOH-CH₂-besteht.
 - 5. Phospholipid-Zusammensetzungen, die durch die folgende allgemeine Formel wiedergegeben werden:

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$$R-Y-O-P-O-\begin{bmatrix}O\\I\\P-O\\A_I\end{bmatrix}_X$$

dadurch gekennzeichnet, dass

- A ausgewählt wird unter H, M oder R-Y-;
- A₁ ausgewählt wird unter H, OH, OM oder R-Y-O-;
- x für 0 oder für eine ganze Zahl von 1 bis 5 steht;
 - M ein Kation ist;

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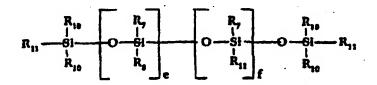
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- Y ein Alkylen oder ein substituiertes Alkylen ist; und
- R eine Mischung aus quaternierten Organosilikon-Aminhälften darstellt, die ausgewählt werden aus der Gruppe bestehend aus:
 - a) einer quaternierten Organosilikon-Amidoaminhälfte mit der Formel:

$$R_{1} = \begin{bmatrix} & & & & & & \\ & & & & \\ & & & & & \\ & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & \\ & & & \\ & & & \\ & & & & \\ & & & \\ & & & & \\ & & & \\ & & & & \\ & & & \\ & & & & \\ & & & \\ & & & \\ & & & & \\ & &$$

in welcher;

R₁ eine Silikonhauptkette ist, an welche eine oder mehrere Funktionsgruppen von Amidoamin oder Amin angehängt werden können und welche wiedergegeben wird durch die Formel:



in welcher,

R₇ und R₈, die identisch oder unterschiedlich sein können, ausgewählt werden unter Alkyl, Aryl, verkapptem oder unverkapptem Polyoxyalkylen, Alkaryl, Aralkylen oder Alkenyl (Vinyl);

R₁₀, identisch oder unterschiedlich sein kann, und ausgewählt wird unter Alkyl, Aryl oder olefinisch (Vinyl) ist;

R₁₁, das identisch oder unterschiedlich sein kann, ausgewählt wird unter R₁₀, -

 $(CH_2)_{n1}$ - B_{n2} - $(CH_2)_{n3}$ -CO- NR_2 - $(CH_2)_n$ - NR_3R_4 , - R_6 - NR_3R_4 - oder einer Mischung derselben, worin R_2 , R_3 , R_4 , R_6 , R_6 , R_6 , R_6 , R_7 , R_8 ,

e eine ganze Zahl von 0 bis 50.000 sein kann;

f eine ganze Zahl von 0 bis 100 sein kann;

R₂ Wasserstoff oder Alkyl, Hydroxyalkyl oder Alkenyl mit bis zu 6 Kohlenstoffatomen ein jedes darstellt, oder Cycloalkyl mit bis zu 6 Kohlenstoffatomen, oder Polyoxyalkylen mit bis zu 10 Kohlenstoffatomen innerhalb der Oxyalkyleneinheit;

R₃ und R₄, die identisch oder unterschiedlich sein können, ausgewählt werden unter Alkyl, Hydroxyalkyl, Carboxyalkyl mit bis zu 6 Kohlenstoffatomen in einem jeden Alkyl, oder Polyoxyalkylen mit bis zu 10 Kohlenstoffatomen; hinzu kommt, dass R₃ und R₄ zusammen genommen werden mit dem Stickstoff, an das sie gebunden sind, und eine N-heterocyclische Verbindung bilden;

X- ein Anion ist;

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n für eine ganze Zahl von 2 bis 12 steht;

n¹ für Null oder für eine ganze Zahl von 1 bis 12 steht;

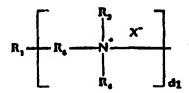
n² für 0 oder für 1 steht;

n³ für eine ganze Zahl von 1 bis 5 steht;

B Schwefel (S) oder Sauerstoff (O) darstellt; unter der Bedingung dass wenn n² gleich 0 ist, n¹ oder n³ mindestens 1 ist, und dass wenn n² gleich 1 ist, n¹ und n³ ein jedes mindestens 1 beträgt;

d für eins oder für mehr steht; und

b) einer quaternierten Organosilikon-tert-Aminhälfte gemäß der Formel:



in welcher,

R₁, wie es oben definiert ist, eine Silikonhauptkette darstellt, an welche Funktionsgruppen von Amidoamin oder Amin angehängt werden können;

R₆, Alkylen, Hydroxyalkylen, Arylen, Alkarylen, Aralkylen, Heteroalkylen darstellt, wobei das Heteroatom N, S oder O sein kann und es kann sich mehr als ein solches Heteroatom in der Kette befinden;

X- ein Anion ist;

d₁ für eine ganze Zahl von 1 oder mehr steht;

R₃ und R₄, die identisch oder unterschiedlich sein können, ausgewählt werden unter Alkyl, Hydroxyalkyl,

Carboxyalkyl mit bis zu 6 Kohlenstoffatomen in einem jeden Alkyl, oder Polyoxyalkylen mit bis zu 10 Kohlenstoffatomen; hinzu kommt, dass $\rm R_3$ und $\rm R_4$ zusammen genommen werden mit dem Stickstoff, an das sie gebunden sind, und eine N-heterocyclische Verbindung bilden;

- und eine quaternierte organische Aminhälfte, die ausgewählt wird aus der Gruppe bestehend aus:
 - c) einer guaternierten organischen Amidoaminhälfte mit der Formel:

$$\begin{bmatrix} O & \begin{bmatrix} R_2 & O \\ \\ R_3 - C & \end{bmatrix} & \begin{bmatrix} R_3 & R_3 \\ \\ R_4 & \end{bmatrix} & \begin{bmatrix} R_3 & C \\ \\ R_4 & \end{bmatrix} & \begin{bmatrix} R_3 & C \\ \\ R_4 & \end{bmatrix}$$

in welcher:

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R₅ Alkyl, Alkenyl, Alkoxyalkyl oder Hydroxyalkyl mit 5 bis 21 Kohlenstoffatomen ein jedes, Alkaryl oder Aryl mit bis zu 20 Kohlenstoffatomen darstellt;

R₂ Wasserstoff oder Alkyl, Hydroxyalkyl oder Alkenyl mit bis zu 6 Kohlenstoffatomen ein jedes darstellt, oder Cycloalkyl mit bis zu 6 Kohlenstoffatomen, oder Polyoxyalkylen mit bis zu 10 Kohlenstoffatomen innerhalb der Oxyalkyleneinheit;

R₃ und R₄, die identisch oder unterschiedlich sein können, ausgewählt werden unter Alkyl, Hydroxyalkyl, Carboxyalkyl mit bis zu 6 Kohlenstoffatomen in einer jeden Alkylhälfte, oder Polyoxyalkylen mit bis zu 10 Kohlenstoffatomen; hinzu kommt, dass R₃ und R₄ zusammen genommen werden mit dem Stickstoff, an das sie gebunden sind, und eine N-heterocyclische Verbindung bilden;

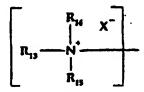
X- ein Anion ist;

g für 0 oder für 1 steht;

n für eine ganze Zahl von 2 bis 12 steht;

n⁴ für 1 oder für mehr steht; und

d). einer organischen quaternierten tert-Aminhälfte mit der Formel:



in welcher

R₁₃, R₁₄ und R₁₅ identisch oder unterschiedlich sind und bestehen aus Alkyl, substituiertem Alkyl, Alkylaryl- oder Alkenylgruppen mit bis zu 16 Kohlenstoffatomen, unter der Bedingung, dass die gesamten Kohlenstoffatome in R₁₃ + R₁₄ + R₁₅ zwischen 10 und 24 betragen;

unter der Bedingung, dass wenn R aus einer Mischung von Organosilikon-Aminhälften und organischen Aminhälften besteht, dann mindestens 5 Äquivalentgewichtsprozente bis zu etwa 60 Äquivalentgewichtsprozente

des gesamten Äquivalenzgewichts der Aminhälften der Phospholipid-Zusammensetzungen eine quaternierte Organosilikon- Amidoaminhälfte, eine quaternierte Organosilikon- tert. - Aminhälfte oder Mischungen derselben darstellen.

- Phospholipid-Zusammensetzungen gemäß Anspruch 5, dadurch gekennzeichnet, dass g in der quaternierten organischen Amidoaminhälfte c) 1 beträgt.
 - 7. Phospholipid-Zusammensetzungen gemäß Anspruch 5, dadurch gekennzeichnet, dass f gleich 0 ist.
- 8. Phospholipid-Zusammensetzungen gemäß Anspruch 5, dadurch gekennzeichnet, dass die Endgruppen R₁₁ durch R₁₀ dargestellt sind und dass f größer als 0 ist.
 - 9. Phospholipid-Zusammensetzungen gemäß Anspruch 5, dadurch gekennzeichnet, dass R eine Mischung ist aus:
- a) quaternierten Organosilikon-Amidoaminhälften mit der Formel:

$$R_{1} = \begin{bmatrix} O & R_{2} & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\ & & & & \\$$

in welcher:

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R₁, wie es in Anspruch 5 definiert ist, eine Silikonhauptkette darstellt, an welche eine oder mehrere Funktionsgruppen von Amidoamin und/oder Amin angehängt werden können;

R₂ Wasserstoff oder Alkyl, Hydroxyalkyl oder Alkenyl mit bis zu 6 Kohlenstoffatomen ein jedes darstellt, oder Cycloalkyl mit bis zu 6 Kohlenstoffatomen, oder Polyoxyalkylen mit bis zu 10 Kohlenstoffatomen innerhalb der Oxyalkyleneinheit;

R₃ und R₄, die identisch oder unterschiedlich sein können, ausgewählt werden unter Alkyl, Hydroxyalkyl, Carboxyalkyl mit bis zu 6 Kohlenstoffatomen in einem jeden Alkyl, oder Polyoxyalkylen mit bis zu 10 Kohlenstoffatomen;

X- ein Anion ist;

n für eine ganze Zahl von 2 bis 12 steht;

n¹ für Null oder für eine ganze Zahl von 1 bis 12 steht;

45 n² für 0 oder für 1 steht;

n³ für eine ganze Zahl von 1 bis 5 steht;

B Schwefel (S) oder Sauerstoff (O) darstellt; unter der Bedingung dass wenn n² gleich 0 ist, n¹ oder n³ mindestens 1 ist, und dass wenn n² gleich 1 ist, n¹ und n³ ein jedes mindestens 1 beträgt;

d für eins oder mehr steht;

c). quaternierte organische Amidoaminhälften mit der Formel:

$$\begin{bmatrix} O & \begin{bmatrix} R_2 & O \\ R_3 & C \end{bmatrix} & \begin{bmatrix} R_2 & R_3 \\ R_4 & C \end{bmatrix} & \begin{bmatrix} R_2 & R_3 \\ R_4 & C \end{bmatrix} & \begin{bmatrix} R_3 & C \\ R_4 & R_4 \end{bmatrix}$$

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in welcher;

 R_5

Alkyl, Alkenyl, Alkoxyalkyl oder Hydroxyalkyl mit 5 bis 21 Kohlenstoffatomen ein jedes, Alkaryl oder Aryl mit bis zu 20 Kohlenstoffatomen darstellt;

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 R_2 Wasserstoff oder Alkyl, Hydroxyalkyl oder Alkenyl mit bis zu 6 Kohlenstoffatomen ein jedes darstellt, oder Cycloalkyl mit bis zu 6 Kohlenstoffatomen, oder Polyoxyalkylen mit bis zu 10 Kohlenstoffatomen innerhalb der Oxyalkyleneinheit;

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R₃ und R₄,

die identisch oder unterschiedlich sein können, ausgewählt werden unter Alkyl, Hydroxyalkyl, Carboxyalkyl mit bis zu 6 Kohlenstoffatomen in einer jeden Alkylhälfte, oder Polyoxyalkylen mit bis zu 10 Kohlenstoffatomen; hinzu kommt, dass R₃ und R₄ zusammen genommen werden mit dem Stickstoff, an das sie gebunden sind, und eine N-heterocyclische Verbindung bilden

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Xein Anion ist;

g

für 0 oder für 1 steht;

n

für eine ganze Zahl von 2 bis 12 steht;

n⁴

für 1 oder für mehr steht; und

unter der Bedingung, dass mindestens 5 Äquivalentgewichtsprozente bis zu etwa 60 Äquivalentgewichtsprozente des gesamten Äquivalenzgewichts der Aminhälften der Phospholipid-Zusammensetzung eine quaternierte Organosilikon -Amidoaminhälfte oder Mischungen derselben darstellen.

10. Verfahren zur Zubereitung neuer Phospholipid-Zusammensetzungen gemäß Anspruch 5, die wiedergegeben werden durch die allgemeine Formel:

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$$R-Y-O-P-O-P-O-A$$

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dadurch gekennzeichnet, dass,

ausgewählt wird unter H, M, oder R-Y-;

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ausgewählt wird unter H, OH, OM oder R-Y-O-

für 0 oder für eine ganze Zahl von 1 bis 5 steht;

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ein Kation ist;

ein Alkylen oder ein substituiertes Alkylen ist; und

R eine Mischung aus quaternierten Amidoaminhälften und/oder tert.-Aminhälften darstellt, wie sie oben definiert sind:

Verfahren das die Reaktion der Kombination eines Reaktionsmittels eines organischen Amidoamins, eines organischen tert.-Amins oder einer Mischung derselben, und eines Reaktionsmittels eines silikonmodifizierten Amidoamins oder eines silikonmodifizierten tert.-Amins oder einer Mischung derselben mit einem Reaktionsmittels eines Polyphosphats, Phosphits oder Phosphatesterhalids in den äquivalenten Gewichtsverhältnissen von etwa 0,7 bis 3,3 der gesamten Amidoamin- oder tert.-Aminreaktionsmittel zu 1 der Polyphosphat-, Phosphit- oder Phosphatesterhalid-Reaktionsmittel aufweist, bis das Aminreaktionsmittel vollständig reagiert hat, unter der Bedingung, dass mindestens 5 Äquivalentgewichtsprozente bis zu etwa 60 Äquivalentgewichtsprozente des gesamten Äquivalenzgewichts der Aminreaktionsmittel aus Silikon bestehen werden, wobei die genannten Polyphosphat-, Phosphitoder Phosphatesterhalidreaktionsmittel der nachfolgenden allgemeinen Formel entsprechen:

in welcher:

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A₂ ausgewählt wird unter H, M und X-Y-;

A₃ ausgewählt wird unter H, OH, OM und X-Y-O-;

x für 0 oder für eine ganze Zahl von 1 bis 5 steht;

M ein Kation ist;

Y ein Alkylen oder ein substituiertes Alkylen ist; und

X ein Halogen ist.

Revendications

1. Compositions de phospholipide de formule

R-Y-O-P-O-P-O-A

caractérisées en ce que

A est choisi parmi H, M et R-Y-;

A₁ est choisi parmi H, OH, OM et R-Y-O-;

x est 0 ou un nombre entier de 1 à 5;

M est un cation;

Y est un alkylène ou un alkylène substitué;

R est un fragment amine d'un organosilicone quaternisé choisi à partir du groupe consistant en:

a) un fragment amidoamine d'organosilicone quaternisé de formule:

$$R_{1} = \begin{bmatrix} CH_{2} & B_{12} & CH_{2} & B_{13} & CH_{2} & B_{14} & CH_{2} & CH_{2} & B_{14} & CH_{2} & CH_{2} & B_{14} & CH_{2} &$$

dans laquelle:

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R₁ est une chaîne d'un squelette de silicone auquel on peut attacher un/des groupe(s) fonctionnel(s) amidoamine(s) et est représenté par la formule:

 $R_{11} - Si - C - S$

dans laquelle:

 $\rm R_7$ et $\rm R_8$, qui peuvent être identiques ou différents sont choisis à partir d'un alkyle, aryle, polyoxyalkylène bloqué ou non bloqué, alkaryle, aralkylène ou alkényle (vinyle);

R₁₀ peut être identique ou différent et est choisi à partir d'un alkyle, aryle ou un hydrocarbure oléfinique (vinyle);

 R_{11} , qui peut être identique ou différent, est choisi parmi R_{10} , - $(CH_2)_{n1}$ - B_{n2} - $(CH_2)_{n3}$ -CO- NR_2 - $(CH_2)_{n-1}$ - NR_3 R₄, -R₆- NR_3 , R₄- ou des mélanges de ceux-ci, dans lesquels R₂, R₃, R₄, R₆, B, n, n¹, n² et n³ sont comme définis ci-dessus; à la condition qu'au moins un parmi R_{11} est une amidoamine;

e peut être un nombre entier de 0 à 50000;

f peut être un nombre entier de 0 à 100;

 $\rm R_2$ est un hydrogène ou un alkyle, hydroxyalkyle ou alkényle jusqu'à 6 atomes de carbone chacun, ou un cycloalkyle jusqu'à 6 atomes de carbone, ou un polyoxyalkylène jusqu'à 10 atomes de carbone à l'intérieur de l'unité oxyalkylène;

 R_3 et R_4 , qui peuvent être identiques ou différents, sont choisis parmi un alkyle, hydroxyalkyle, carboxyalkyle jusqu'à 6 atomes de carbone dans chaque alkyle, ou polyoxyalkylène jusqu'à 10 atomes de carbone; X- est un anion, de préférence un halogène;

n est un nombre entier de 2 à 12;

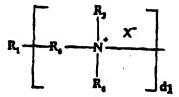
n¹ est zéro ou un nombre entier-de 1 à 12;

n² est 0 ou 1;

n³ est un nombre entier de 1 à 5;

B est du soufre (S) ou un oxygène (O); à la condition que lorsque n² est 0, n¹ ou n³ est au moins 1 et lorsque n² est 1, n¹ et n³ chacun est au moins 1; et d est un ou supérieur;

b) un fragment d'une amine tertiaire d'un organosilicone quaternisé de formule:



dans laquelle:

R₁ tel que défini ci-dessus est une chaîne d'un squelette de silicone auquel des groupes fonctionnels amine ou amidoamine peuvent être attachés;

 R_6 est un alkylène, un hydroxyalkylène, un arylène, un alkarylène, un aralkylène, un hétéroalkylène dans lequel l'hétéroatome peut être N, S ou O et il peut y avoir plus d'un tel hétéroatome dans la chaîne;

X⁻ est un anion;

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d1 est un nombre entier de un ou supérieur;

R₃ et R₄ sont tels que définis ci-dessus;

et des mélanges de ceux-ci.

- 2. Compositions de phospholipides selon la revendication 1, caractérisées en ce que f est 0.
 - Compositions de phospholipides selon la revendication 1, caractérisées en ce que les groupes terminaux R₁₁ sont R₁₀ et f est supérieur à 0.
- 4. Compositions de phospholipides selon la revendication 1, caractérisées en ce que Y est -CH₂-CHOH-CH₂-.
 - 5. Compositions de phospholipides représentées par la formule générale suivante:

R-Y-O-P-O-P-O-A

caractérisées en ce que

A est choisi parmi H, M, ou R-Y-;

A₁ est choisi parmi H, OH, OM ou R-Y-O-;

x est 0 ou un nombre entier de 1 à 5;

M est un cation;

Y est un alkylène ou un alkylène substitué; et

R est un mélange de fragment amine d'organosilicone quaternisé choisi à partir du groupe consistant en:

a) un fragment amidoamine d'organosilicone quaternisé de formule:

 $R_{1} = \begin{bmatrix} & & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & & \\ & & & & \\ & & & & \\ & & & & \\ &$

dans laquelle:

R₁ est une chaîne d'un squelette de silicone auquel on peut attacher des groupes fonctionnels amines ou amidoamines et est représenté par la formule:

$$R_{11} - \begin{array}{c|c} R_{10} & R_{7} & R_{7} \\ \hline R_{10} & Si & O - Si \\ R_{10} & R_{0} & R_{11} \end{array}$$

dans laquelle:

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R₇ et R₈, qui peuvent être identiques ou différents, sont choisis à partir d'un alkyle, aryle, polyoxyalkylène bloqué ou non bloqué, alkaryle, aralkylène ou alkényle (vinyle);

R₁₀ peut être identique ou différent et est choisi parmi d'un alkyle, aryle ou d'un hydrocarbure oléfinique (vinyle) ;

 R_{11} , qui peut identique ou différent, est choisi parmi R_{10} , - $(CH_2)_{n1}$ - B_{n2} - $(CH_2)_{n3}$ -CO- NR_2 - $(CH_2)_n$ - NR_3R_4 , - R_6 - NR_3 - R_4 - ou des mélanges de ceux-ci, dans lequel R_2 , R_3 , R_4 , R_6 , R_6 , R_7 , R_8 ,

e peut être un nombre entier de 0 à 50000;"

f peut être un nombre entier de 0 à 100;

 $\rm H_2$ est un hydrogène ou un alkyle, hydroalkyle ou alkényle jusqu'à 6 atomes de carbone chacun, ou un cycloalkyle jusqu'à 6 atomes de carbone, ou un polyoxyalkylène jusqu'à 10 atomes de carbone à l'intérieur de l'unité oxyalkylène;

R₃ et R₄, qui peuvent être identiques ou différents, sont choisis à partir d'un alkyle, hydroxyalkyle, carboxyalkyle jusqu'à 6 atomes de carbone dans chaque alkyle, ou un polyoxyalkylène jusqu'à 10 atomes de carbone; de plus R₃ et R₄, pris ensemble avec l'azote auquel ils sont attachés, représentent un N-hétérocycle;

X- est un anion;

n est un nombre entier de 2 à 12;

n1 est zéro ou un nombre entier de 1 à 12;

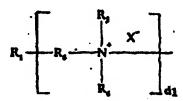
n² est 0 ou 1:

n3 est un nombre entier de 1 à 5;

B est un soufre (S) ou un oxygène (O); à la condition que lorsque n² est 0, n¹ ou n³ est au moins 1 et lorsque n² est 1, n¹ et n³ chacun est au moins 1; et

d est un ou supérieur; et

b) un fragment amine tertiaire d'un organosilicone quaternisé de formule:



dans laquelle:

R₁ tel que défini ci-dessus est une chaîne d'un squelette de silicone à laquelle on peut attacher des groupes fonctionnels amidoamine ou amine peuvent être attachés;

R₆ est un alkylène, hydroxyalkylène, arylène, alkarylène, aralkylène, hétéroalkylène dans lequel l'hétéromatome peut être N, S ou O et il peut y avoir plus d'un tel hétéroatome dans la chaîne;

X⁻ est un anion;

d1 est un nombre entier de un ou supérieur;

R₃ et R₄, qui peuvent être identiques ou différents, sont choisis à partir d'un alkyle, hydroxyalkyle, carboxyalkyle jusqu'à 6 atomes de carbone dans chaque alkyle, ou polyoxyalkylène jusqu'à 10 atomes de carbone; de plus R₃ et R₄ pris ensemble avec l'azote auquel ils sont attachés représentent un N-hétérocycle.

et un fragment d'une amine organique quaternisée choisi à partir du groupe consistant en:

c) un fragment d'une amidoamine organique quaternisée de formule:

$$\begin{bmatrix} O & \begin{bmatrix} R_2 & O \\ \\ R_3 - C & \end{bmatrix} & \begin{bmatrix} R_2 & R_3 \\ \\ R_4 & \end{bmatrix} & \begin{bmatrix} R_3 & X^- \\ \\ R_4 & \end{bmatrix} \end{bmatrix}$$

dans laquelle:

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 $\rm R_{5}$ est un alkyle, alkényle, alkoxyalkyle ou hydroxyalkyle de 5 à 21 atomes de carbone chacun, alkaryle ou aryle jusqu'à 20 atomes de carbone;

 $\rm R_2$ est un hydrogène ou alkyle, hydroxyalkyle ou alkényle jusqu'à 6 atomes de carbone chacun, cycloalkyle jusqu'à 6 atomes de carbone, ou polyoxyalkylène jusqu'à 10 atomes de carbone à l'intérieur de l'unité oxyalkylène;

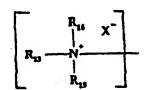
 R_3 et R_4 , qui peuvent être identiques ou différents, sont choisis à partir d'un alkyle, hydroalkyle, carboxyalkyle jusqu'à 6 atomes de carbone dans chaque fragment alkyle, et polyoxyalkylène jusqu'à 10 atomes de carbone; de plus R_3 et R_4 pris ensemble avec l'azote auquel ils sont attachés représentent un hétérocycle; X^* est un anion;

g est 0 ou 1;

n est un nombre entier de 2 à 12; et

n4 est 1 ou supérieur; et

d) un fragment d'une amine organique tertiaire quaternisée de formule:



dans laquelle:

 R_{13} , R_{14} et R_{15} sont identiques ou différents et sont des groupes alkyles, alkyles substitués, alkylaryles ou alkényles jusqu'à 16 atomes de carbone à la condition que le total des atomes de carbone dans R_{13} + R_{14} + R_{15} est entre 10 et 24;

à la condition que lorsque R est un mélange de fragment d'amine organique et d'amines d'organosilicones, au moins 5 équivalents % en poids jusqu'à environ 60 équivalents % en poids de l'équivalent en poids total des fragments amines de la composition de phospholipide est un fragment amidoamine d'organosilicone quaternisé, un fragment amine tertiaire d'organosilicone quatemisé ou des mélanges de ceux-ci.

- 6. Compositions de phospholipides selon la revendication 5, caractérisées en ce que g dans le fragment amidoamine organique quaternisé c) est 1.
- 7. Compositions de phospholipides selon la revendication 5, caractérisées en ce que f est 0.
- Compositions de phospholipides selon la revendication 5, caractérisées en ce que les groupes terminaux R₁₁ sont R₁₀ et f est supérieur à 0.
- 9. Compositions de phospholipides selon la revendication 5, caractérisées en ce que R est un mélange de:
 - a) fragments amidoamine d'organosilicone quaternisé de formule:

$$R_{1} = \begin{bmatrix} O & R_{2} & & & \\$$

dans laquelle:

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 R_1 comme défini dans la revendication 5 est une chaîne d'un squelette de silicone auquel on peut attacher des groupes fonctionnels amidoamine et/ou amine;

R₂ est un hydrogène ou un alkyle, hydroxyalkyle ou alkényle jusqu'à 6 atomes de carbone chacun, ou cycloalkyle jusqu'à 6 atomes de carbone, ou polyoxyalkylène jusqu'à 10 atomes de carbone à l'intérieur de l'unité oxyalkylène;

R₃ et R₄, qui peuvent être identiques ou différents, sont choisis à partir d'un alkyle, hydroxyalkyle, carboxyalkyle jusqu'à 6 atomes de carbone dans chaque alkyle, ou un polyoxyalkylène jusqu'à 10 atomes de carbone:

X- est un anion:

n est un nombre entier de 2 à 12;

n¹ est zéro ou un nombre entier de 1 à 12;

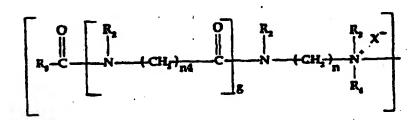
n² est 0 ou 1:

n³ est un nombre entier de 1 à 5;

B est un soufre (S) ou un oxygène (O); à la condition que lorsque n² est 0, n¹ ou n³ est au moins 1 et lorsque n² est 1, n¹ et n³ chacun est au moins 1; et

d est un ou supérieur; et

c) les fragments amidoamine organique quaternisé de formule:



dans laquelle:

 $\rm R_5$ est.un alkyle, alkényle, alkoxy, alkyle ou hydroxyalkyle de 5 à 21 atomes de carbone chacun, alkaryle ou aryle jusqu'à 20 atomes de carbone;

R₂ est un hydrogène ou un alkyle, hydroxyalkyle ou alkényle jusqu'à 6 atomes de carbone chacun, cycloalkyle jusqu'à 6 atomes de carbone, ou polyoxyalkylène jusqu'à 10 atomes de carbone à l'intérieur de l'unité oxyalkylène;

R₃ et R₄, qui peuvent être identiques ou différents, sont choisis à partir d'un alkyle, hydroxyalkyle, carboxyalkyle jusqu'à 6 atomes de carbone dans chaque fragment alkyle, ou polyoxyalkylène jusqu'à 10 atomes de carbone; de plus R₃ et R₄ pris ensemble avec l'azote auquel ils sont attachés présentent un N-hétérocycle;

X est un anion;

g est 0 ou 1;

n est un nombre entier de 2 à 12;

n4 est 1 ou supérieur;

à la condition qu'au moins 5 équivalents % en poids jusqu'à environ 60 équivalents % en poids des équivalents en poids total des fragments amines de la composition de phospholipides sont un fragment amidoamine d'un organosilicone quaternisé ou des mélanges de ceux-ci.

10. Méthode de préparation de nouvelles compositions de phospholipides selon la revendication 5 représentées par la formule générale:

caractérisée en ce que

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A est choisi parmi H, M ou R-Y-;
A₁ est choisi parmi H, OH, OM ou R-Y-O-;
x est 0 ou un nombre entier de 1 à 5;
M est un cation:

Y est un alkylène ou un alkylène substitué; et

R est un mélange d'une amidoamine quaternisée et/ou de fragments d'amine tertiaire tels que définis cidessus;

qui comprend de faire réagir la combinaison d'un réactif d'une amine tertiaire organique d'amidoamine organique ou un mélange de celui-ci et une amidoamine modifiée par un silicone ou un réactif d'amine tertiaire modifiée par un silicone ou des mélanges de ceux-ci avec un réactif d'un halogénure de phosphate ester, de phosphite ester ou de polyphosphate ester dans des rapports en poids équivalents d'environ 0,7 à 3,3, des réactifs d'amine tertiaire ou d'amidoamine totale pour 1 réactif d'halogénure de phosphate ester, de phosphite ester ou de polyphosphate jusqu'à ce que le réactif amine ait complètement réagi, à la condition qu'au moins 5 équivalents % en poids jusqu'à 60 équivalents % en poids équivalent total des réactifs amines contienne du silicone, ce réactif d'halogénure de phosphate ester, de phosphite ester ou de polyphosphate ester ayant la formule générale:

dans laquelle:

A₂ est choisi parmi H, M et X-Y-; A₃ est choisi parmi H, OH, OM et X-Y-O-; x est 0 ou un nombre entier de 1 à 5; M est un cation; Y est un alkylène ou un alkylène substitué; et X est un halogène.

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